

Claims

1. Nutritional and pharmaceutical formulations comprising in combination a source of vitamin K and a source of at least one essential fatty acid (EFA), in which the concentration of vitamin K is not less than 1000 $\mu\text{g}/100\text{g}$.
2. Nutritional and pharmaceutical formulations according to claim 1, in which the concentration of vitamin K is not less than 1000 $\mu\text{g}/10\text{g}$.
3. Nutritional and pharmaceutical formulations according to claim 1 which provide a daily dose between 50 μg and 100 mg vitamin K and between 50 mg and 100 g of the EFA.
4. Nutritional and pharmaceutical formulations according to claim 1 in which the form of vitamin K used is phylloquinone (vitamin K1).
5. Nutritional and pharmaceutical formulations according to claim 1 in which the EFA is selected from gamma-linolenic acid, dihomogammalinolenic acid, arachidonic acid and adrenic acid, and combinations of these EFAs.
6. Nutritional and pharmaceutical formulations according to claim 1 in which the EFA is selected from stearidonic acid, eicosapentaenoic acid, docosapentaenoic acid and docosahexenoic acid, and combinations of these EFAs.

7. Nutritional and pharmaceutical formulations according to claim 1 in which there is at least one n-6 EFA and at least one n-3 EFA present, the n-6 EFA(s) selected from gamma-linolenic acid, dihomogammalinolenic acid, arachidonic acid and adrenic acid, and combinations of these acids, and the n-3 EFA(s) selected from stearidonic acid, eicosapentaenoic acid, docosapentaenoic acid and docosahexaenoic acid, and combinations of these acids.
8. Nutritional and pharmaceutical formulations according to claim 1 in which the active ingredient consists essentially wholly of EFA and vitamin K.
9. Nutritional and pharmaceutical formulations according to claim 1 further comprising one or more essential vitamins and/or minerals or one or more pharmaceutical drugs.
10. Nutritional and pharmaceutical formulations comprising in combination a source of vitamin K and a source of at least one essential fatty acid (EFA), in which proteins and amino acids are absent from the active ingredients of the formulation.
11. Nutritional and pharmaceutical formulations according to claim 10 which provide a daily dose between 50 μ g and 100 mg vitamin K and between 50 mg and 100 g of the EFA.

12. Nutritional and pharmaceutical formulations according to claim 10 in which the form of vitamin K used is phylloquinone (vitamin K1).
- 5 13. Nutritional and pharmaceutical formulations according to claim 10 in which the EFA is selected from gamma-linolenic acid, dihomogammalinolenic acid, arachidonic acid and adrenic acid, and combinations of these EFAs.
- 10 14. Nutritional and pharmaceutical formulations according to claim 10 in which the EFA is selected from stearidonic acid, eicosapentaenoic acid, docosapentaenoic acid and docosahexenoic acid, and combinations of these EFAs.
- 15 15. Nutritional and pharmaceutical formulations according to claim 10 in which there is at least one n-6 EFA and at least one n-3 EFA present, the n-6 EFA(s) selected from gamma-linolenic acid, dihomogammalinolenic acid, arachidonic acid and adrenic acid, and combinations of these
20 acids, and the n-3 EFA(s) selected from stearidonic acid, eicosapentaenoic acid, docosapentaenoic acid and docosahexaenoic acid, and combinations of these acids.
- 25 16. Nutritional and pharmaceutical formulations according to claim 10 in which the active ingredient consists essentially wholly of EFA and vitamin K.
17. Nutritional and pharmaceutical formulations according to claim 10 further comprising one or

stress, mental, psychological, psychiatric
or neurological disorders;

skin disorders;

asthma or other respiratory disorder;

arthritis or any form of inflammatory,
gastrointestinal, kidney or reproductive system
disorder;

25. The method according to claim 22 in which the form of vitamin K used is phylloquinone (vitamin K1).

26. The method according to claim 22 in which the EFA is selected from gamma-linolenic acid, dihomo-gammalinolenic acid, arachidonic acid and adrenic acid, and combinations of these EFAs.

eicosapentaenoic acid, docosapentaenoic acid and docosahexenoic acid, and combinations of these EFAs.

28. The method according to claim 22 in which there
5 is at least one n-6 EFA and at least one n-3 EFA present, the n-6 EFA(s) selected from gamma-linolenic acid, dihomogammalinolenic acid, arachidonic acid and adrenic acid, and combinations of these acids, and the n-3 EFA(s)
10 selected from stearidonic acid, eicosapentaenoic acid, docosapentaenoic acid and docosahexaenoic acid, and combinations of these acids.
29. The method according to claim 22 in which the
15 active ingredient consists essentially wholly of EFA and vitamin K.
30. The method according to claim 22 further comprising one or more essential vitamins and/or minerals or one or more pharmaceutical drugs.
- 20 31. A method of treating or preventing a variety of diseases or conditions including:

premenstrual or menstrual disorders of any kind;

- 25 bone or calcium disorders of any kind, including osteoporosis;

metabolic or cardiovascular disorders including diabetes, obesity, elevated blood cholesterol or triglyceride levels or cardiovascular disorders;

stress, mental, psychological, psychiatric
or neurological disorders;

skin disorders;

asthma or other respiratory disorder;

5 arthritis or any form of inflammatory,
gastrointestinal, kidney or reproductive system
disorder;

10 using nutritional and pharmaceutical formulations
comprising in combination a source of vitamin K
and a source of at least one essential fatty acid
(EFA), in which proteins and amino acids are
absent from the active ingredients of the
formulation.

15 32. The method according to claim 31 which provides a
daily dose between 50 μ g and 100 mg vitamin K and
between 50 mg and 100 g of the EFA.

33. The method according to claim 31 in which the
form of vitamin K used is phylloquinone (vitamin
K1).

20 34. The method according to claim 31 in which the EFA
is selected from gamma-linolenic acid,
dihomogammalinolenic acid, arachidonic acid and
adrenic acid, and combinations of these EFAs.

25 35. The method according to claim 31 in which the EFA
is selected from stearidonic acid,
eicosapentaenoic acid, docosapentaenoic acid and
docosahexenoic acid, and combinations of these
EFAs.

36. The method according to claim 31 in which there is at least one n-6 EFA and at least one n-3 EFA present, the n-6 EFA(s) selected from gamma-linolenic acid, dihomogammalinolenic acid,
5 arachidonic acid and adrenic acid, and combinations of these acids, and the n-3 EFA(s) selected from stearidonic acid, eicosapentaenoic acid, docosapentaenoic acid and docosahexaenoic acid, and combinations of these acids.
- 10 37. The method according to claim 31 in which the active ingredient consists essentially wholly of EFA and vitamin K.
38. The method according to claim 31 further comprising one or more essential vitamins and/or
15 minerals or one or more pharmaceutical drugs.
39. A method of treating or preventing a variety of diseases or conditions including:
- premenstrual or menstrual disorders of any kind;
 - 20 bone or calcium disorders of any kind, including osteoporosis;
 - metabolic or cardiovascular disorders including diabetes, obesity, elevated blood cholesterol or triglyceride levels or
25 cardiovascular disorders;
 - stress, mental, psychological, psychiatric or neurological disorders;
 - skin disorders;
 - asthma or other respiratory disorder;

arthritis or any form of inflammatory,
gastrointestinal, kidney or reproductive system
disorder;

5 using foodstuff which already contain EFAs to
which have been added vitamin K in an amount to
raise the vitamin K content of the food to 1000
 μg / 100 g food, or more.

40. The method according to claim 39 using foodstuff
which already contain EFAs to which have been
10 added vitamin K in an amount to raise the vitamin
K content of the food to 1000 μg / 10 g food, or
more.

41. The method according to claim 39 in which the
specific EFA(s) content has been raised by the
15 addition of one of more EFAs.

42. The method according to claim 39 using foodstuff
which naturally contains clinically or
nutritionally small amounts of vitamin K and / or
EFA(s) to which has been added vitamin K and
20 EFAs.